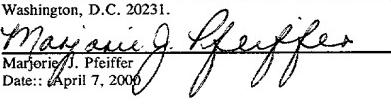


FORM PTO-1390U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE (REV 5-93)		ATTORNEY'S DOCKET NUMBER
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		9052-53
INTERNATIONAL APPLICATION NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED
PCT/GB98/03073	12 October 1998	10 October 1997 ✓
TITLE OF INVENTION		
<i>ELECTROMAGNETIC RADIATION THERAPY</i> ✓		
APPLICANT(S) FOR DO/EO/US		
Gordon Rex Paterson DOUGAL ✓		
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:		
<ol style="list-style-type: none"> <li>1. <input checked="" type="checkbox"/> This is a <b>FIRST</b> submission of items concerning a filing under 35 U.S.C. 371.</li> <li>2. <input type="checkbox"/> This is a <b>SECOND</b> or <b>SUBSEQUENT</b> submission of items concerning a filing under 35 U.S.C. 371.</li> <li>3. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(l).</li> <li>4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.</li> <li>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))             <ol style="list-style-type: none"> <li>a. <input checked="" type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau).</li> <li>b. <input type="checkbox"/> has been transmitted by the International Bureau.</li> <li>c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</li> </ol> </li> <li>6. <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)).</li> <li>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))             <ol style="list-style-type: none"> <li>a. <input checked="" type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau).</li> <li>b. <input type="checkbox"/> have been transmitted by the International Bureau.</li> <li>c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</li> <li>d. <input type="checkbox"/> have not been made and will not be made.</li> </ol> </li> <li>8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</li> <li>9. <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).</li> <li>10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</li> </ol>		
Items 11. to 16. below concern other document(s) or information included:		
<ol style="list-style-type: none"> <li>11. <input checked="" type="checkbox"/> An Information Disclosure Statement under 37 C.F.R. 1.97 and 1.98.</li> <li>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</li> <li>13. <input checked="" type="checkbox"/> A <b>FIRST</b> preliminary amendment. <input type="checkbox"/> A <b>SECOND</b> or <b>SUBSEQUENT</b> preliminary amendment.</li> <li>14. <input type="checkbox"/> A substitute specification.</li> <li>15. <input type="checkbox"/> A change of power of attorney and/or address letter.</li> <li>16. <input checked="" type="checkbox"/> Other items or information: PCT Request; International Search Report; International Preliminary Examination Report</li> </ol>		

U.S. APPLICATION NO. (If known, see 37 C.F.R. 1.50) <b>09/529210</b>	INTERNATIONAL APPLICATION NO. PCT/GB98/03073	ATTORNEY'S DOCKET NUMBER 9052-53	
17. [X] The following fees are submitted:		CALCULATIONS	PTO USE ONLY
<b>Basic National Fee (37 CFR 1.492(a)(1)-(5)):</b> Search Report has been prepared by the EPO or JPO . . . . . \$840.00			
International preliminary examination fee paid to USPTO (37 CFR 1.482) . . . . . \$670.00			
No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)) . . . . . \$760.00			
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO. . . . . \$970.00			
International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4). . . . . \$96.00			
ENTER APPROPRIATE BASIC FEE AMOUNT = \$840.00		\$ 840.00	
Surcharge of <b>\$130.00</b> for furnishing the oath or declaration later than [ ] 20 [ ] 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		\$	
Claims	Number Filed	Number Extra	Rate
Total Claims	30-20 =	10	X \$18.00
Independent Claims	3-3 =	0	X \$78.00
Multiple dependent claim(s) (if applicable)		+ \$260.00	\$
TOTAL OF ABOVE CALCULATIONS =			\$1,020.00
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).		\$	
SUBTOTAL =		\$1,020.00	
Processing fee of <b>\$130.00</b> for furnishing the English translation later than [ ] 20 [ ] 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$	
TOTAL NATIONAL FEE =		\$1,020.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). <b>\$40.00</b> per property +		\$	
TOTAL FEES ENCLOSED =		\$1,020.00	
		Amount to be refunded	\$
		charged	\$
a. [X] A check in the amount of <b>\$1,020.00</b> to cover the above fees is enclosed.			
b. [ ] Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.			
c. [X] The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 50-0220.			
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.			
SEND ALL CORRESPONDENCE TO:		"Express Mail" mailing label number EL533612790US Date of Deposit: April 7, 2000	
Robert J. Smith Myers Bigel Sibley & Sajovec Post Office Box 37428 Raleigh, North Carolina 27627		I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to BOX PCT, Assistant Commissioner for Patents, Washington, D.C. 20231.  Marjorie J. Pfeiffer Date: April 7, 2000	
 SIGNATURE			
40,820 REGISTRATION NUMBER			

Attorney's Docket No. 9052-53

PATENT

IN THE UNITED STATES DESIGNATED OFFICE (DO/US)

In re: Application of Dougal  
Serial No.: To be Assigned  
Filed: Concurrently Herewith  
For: ELECTROMAGNETIC RADIATION  
THERAPY

Date: April 7, 2000

BOX PCT  
Assistant Commissioner for Patents  
Washington, DC 20231

PRELIMINARY AMENDMENT

Sir:

Prior to the examination of the above application, please amend the above-identified application as follows:

In the Claims:

2. (Amended) A electromagnetic radiation therapy system according to Claim 1 [either of Claims 1 or 2] wherein the wavelength is in the range 980 nm-1300 nm.

3. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the wavelength is at, or about, 1072 nm.

4. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the wavelength is at, or about, 1268 nm.

5. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the half angle divergence of the electromagnetic radiation is in the range 15° to 45°.

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Serial No.: To be assigned  
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6. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the electromagnetic radiation is continuous or pulsed.

7. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any preceding claim] wherein, in the instance of the electromagnetic radiation being continuous, the intensity is at least 50  $\mu\text{Watts}/\text{cm}^2$  for the treatment of eyes and mucous membranes and up to 2  $\text{Watts}/\text{cm}^2$ .

8. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any preceding claim] wherein, in the instance of the electromagnetic radiation being continuous, the intensity is at least 500  $\mu\text{Watts}/\text{cm}^2$  for the treatment of skin and up to 2  $\text{Watts}/\text{cm}^2$ .

9. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-6] wherein, in the instance of the electromagnetic radiation being pulsed, the intensity is at least 50  $\mu\text{Watts}/\text{cm}^2$  peak power for the treatment of eyes and mucous membranes and the average power is up to 2  $\text{Watts}/\text{cm}^2$ .

10. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-6] wherein, in the instance of the electromagnetic radiation being pulsed, the intensity is at least 500  $\mu\text{Watts}/\text{cm}^2$  peak power for the treatment of skin and the average power is up to 2  $\text{Watts}/\text{cm}^2$ .

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11. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-6 or 9 or 10] wherein the average power of the pulsed electromagnetic radiation intensity is in the region of 50-100  $\mu$ Watts/cm<sup>2</sup>.

12. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-7 or 9-11] wherein the pulsed electromagnetic radiation is applied for periods of at least 10-15 microseconds.

13. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-7 or 9-12] wherein the pulsed electromagnetic radiation is applied at a frequency/repetition rate in the range of 480-800 Hz.

15. (Amended) An electromagnetic radiation therapy system according to Claim 6 [any of Claims 1-7 or 9-14] wherein the pulsed electromagnetic radiation is applied to the affected area for at least 30 seconds and up to 15 minutes.

16. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the electromagnetic radiation therapy system also includes means for reducing the amount of ambient radiation which impinges on the site of treatment.

18. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] further including means for fixing the intensity of the radiation within a predetermined range.

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Serial No.: To be assigned  
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Page 4 of 5

19. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein radiation output is monitored with a visible display indicating correct function of the device both for intensity and wavelength.

20. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim further including] further including means for controlling the duration of the application of the radiation.

21. (Amended) An electromagnetic radiation therapy system according to Claim 1 [any preceding claim] wherein the radiation producing means are solid state light emitting devices.

23. (Amended) An electromagnetic radiation therapy system according to [either] Claim 21 [or 22] wherein radiation from [such] said solid state light emitting devices is electrically operated or delivered to an applicator via a fibre-optic delivery system.

24. (Amended) An electromagnetic radiation therapy system according to Claim 21 [any of Claims 21-23] wherein the radiation emitter includes a PN junction arranged to emit radiation with a wavelength centring at, or about, 1072 nm or at, about, 1268 nm.

In re: Application of Dougal  
Serial No.: To be assigned  
Filed: Concurrently herewith  
Page 5 of 5

## REMARKS

Claims 1-30 are presented for examination. The above claims have been amended to better conform to U.S. practice. Applicants respectfully request substantive examination on the merits.

Respectfully submitted,



Robert J. Smith  
Registration No. 40,820

Correspondence Address:  
USPTO Customer No.: **20792**  
Myers Bigel Sibley & Sajovec  
Post Office Box 37428  
Raleigh, NC 27627  
Telephone (919) 854-1400  
Facsimile (919) 854-1401

"Express Mail" mailing label number EL533612790US  
Date of Deposit: April 7, 2000

I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to BOX PCT, Assistant Commissioner for Patents, Washington, DC 20231.

  
Marjorie J. Pfeiffer  
Date of Signature: April 7, 2000

ELECTROMAGNETIC RADIATION THERAPY

**Field of the Invention**

- 5 This invention relates to an apparatus producing, and a method of therapy using, electromagnetic radiation for the treatment of diseases and for the maintenance or improvement of organs or body tissues, including muscles. The invention may be used in connection with the cure or alleviation of a variety of diseases including infectious diseases and pathological processes including those caused by viruses and  
10 bacteria.

By way of example, the invention may be used in connection with diseases caused by the herpes virus which is known to be responsible for a number of common ailments including corneal dendritic ulcers, genital herpes, herpes labialis (cold sores), herpes  
15 zoster (shingles) and herpes stomatitis. These infections tend to be recurrent and are not cured by existing, medically accepted treatments.

**Background of the Invention**

- 20 Current medically accepted methods of treating infections caused by the herpes virus are chemotherapeutic agents which are applied topically, injected or taken orally. Such treatment can often deal with the immediate infection but does not prevent a recurrence of the infection at a later date after the treatment has ceased.
- 25 It has been known for several decades that the use of light can give a positive therapeutic effect in the treatment of a wide spectrum of diseases. In the 1960's the use of narrow wavelength light was investigated in *in vivo/in vitro* experiments. It was found that light of wavelength greater than 440nm did not work. Further investigations were carried out with light having a wavelength of from 300 to 350nm  
30 (UV light) but it was found that infection was exacerbated/promoted rather than ameliorated/eliminated. Some attempts have been made to treat individuals affected

with the herpes virus by treatment with light of the wavelength 660nm, as described in US 5500009. However, the present inventor was unable to achieve a significant clinical outcome or benefit at that wavelength.

- 5    Additionally, it is known from the prior art to use a laser to produce coherent radiation and to focus it on the area to be treated. Nd YAG laser treatment at a fundamental wavelength of 1064 nm is associated with decreased pain, scarring and improved healing (US 5445146). Additionally it has been reported that diodes emitting light at the red wavelength,  $940 \pm 25$  nm can be used to treat a range of  
10    essentially musculoskeletal ailments (US 5259380). However there is no indication that light of a wavelength above this would be of any therapeutic use.

- 15    It has now been surprisingly established that low intensity electromagnetic radiation of small bandwidth is effective in the treatment of infectious diseases, inflammatory-type diseases and other conditions, including the alleviation of pain. It is postulated that the way in which the electromagnetic radiation effects its action is by way of energy transmission through cellular components/organelles.

- 20    A water molecule that has a range of electromagnetic radiation wavelengths passed through it will produce several transmission peaks. These transmission peaks are associated with the preferred therapeutic electromagnetic radiation wavelength range of the invention and thus implies a role for the water molecule in the general mechanism of action.

25    **Statements of the Invention**

- According to the present invention there is provided an electromagnetic radiation therapy system comprising means for emitting divergent electromagnetic radiation between 980 and 1500 nm and being capable of producing, at the site being treated, a  
30    radiation intensity of at least  $50 \mu\text{Watts}/\text{cm}^2$  and up to  $2 \text{ Watts}/\text{cm}^2$ .

Reference herein to a site being treated is intended to include, without limitation, the skin or musculature or internal organ of a human or animal subject.

- Preferably the wavelength of the electromagnetic radiation is in the range 980nm-  
5 1300nm. A particularly preferred wavelength is at, or about, 1072nm. A yet further particularly preferred wavelength is at, or about, 1268nm.

- Our studies have shown that the wavelength centred around 1072 nm is particularly effective at treating herpetic and bacterial infections, alleviating acute pain and in  
10 treating eye conditions, whilst the wavelength centred around 1268 nm is particularly effective at providing pain relief from deep muscle injury. It is of note that these two preferred wavelengths correspond to the peak emission wavelengths of a water molecule light transmission profile and thus we believe that the mechanism of action is related to water and possibly cell membranes.

- 15 By divergent it meant that the electromagnetic radiation emitted from the system of the invention has a divergent half angle of at least 5°. Preferably divergence of the electromagnetic radiation is in the range 15° to 45° half angled divergent.

- 20 Preferably the electromagnetic radiation is continuous or pulsed.

Preferably when the electromagnetic radiation is continuous the intensity is at least 50  $\mu$ Watts/cm<sup>2</sup> for treatment of eyes and mucous membranes, and more preferably is at least 500  $\mu$ Watts/cm<sup>2</sup> for treatment of skin and up to 2 Watts/cm<sup>2</sup>.

- 25 Preferably when the electromagnetic radiation is pulsed the intensity is at least 50  $\mu$ Watts/cm<sup>2</sup> peak power for treatment of eyes and mucous membranes, and more preferably is at least 500  $\mu$ Watts/cm<sup>2</sup> peak power for treatment of skin and the average power is up to 2 Watts/cm<sup>2</sup>. The average power is the peak power multiplied  
30 by the proportion of the total time that the radiation is applied. For instance if the

peak power is 500  $\mu$ Watts/cm<sup>2</sup> and is pulsed for 10  $\mu$ seconds at a frequency of 600 Hz then the average power is 30 $\mu$  Watts/ cm<sup>2</sup>.

5 Preferably when the electromagnetic radiation is pulsed the average power of the intensity is in the region of 50-100 $\mu$  Watts/ cm<sup>2</sup>.

We have found that the power may suitably range from 500  $\mu$ Watts/cm<sup>2</sup> peak to 2 Watts/cm<sup>2</sup> continuous or peak power when applied to the skin. In the instance of applying electromagnetic radiation therapy to the eye or mucous membrane, powers 10 as low as 50 $\mu$ Watts/cm<sup>2</sup> continuous or pulsed are found to be beneficial. Typically 10 mWatts/cm<sup>2</sup> are used on skin but this value is dependent on how fat or muscular the subject is and thus how deep the tissue/area/organ to be treated may lie beneath the skin surface. Typically radiation of the intensity 5 mWatts/cm<sup>2</sup> is used on mucous membranes.

15

Preferably when the electromagnetic radiation is pulsed it is applied for periods of at least 10-15  $\mu$ seconds and more preferably is applied at a frequency/repetition rate in the range 480-800 Hz more preferably still the frequency/repetition rate is at, or about, 600 Hz.

20

Our studies have shown that the electromagnetic radiation can be either coherent or non-coherent the clinical outcomes are not affected by this parameter.

25

Preferably the electromagnetic radiation is applied to the affected area for at least 30 seconds and upto a few minutes. A typical exposure time for the skin or eye is in the region of 3 minutes, however for tissues well below the skin surface this time is increased according to the individuals fat/muscle layer depth and exposure could be up to 10 minutes.

30

It should be appreciated that the power source emitting the electromagnetic radiation will have to produce more than the required intensity for the clinical effect since we

have shown that approximately 99% of the applied therapeutic amount of light is lost across the skin surface during treatment. Thus the intensity of applied radiation will have to be corrected for when carrying out a treatment.

- 5 Our studies have shown that the first clinical effects can be detected following 30 seconds of treatment for herpetic infections and that the majority of immediate clinical effects are experienced following 90 seconds of treatment. However certain tissues are more sensitive, for example the mucous membranes are exposed to the dosing regimen for approximately 30 seconds and immediate clinical effects are  
10 noted after only 3 seconds of treatment.

From the foregoing it is understood that the electromagnetic radiation may be directed to the target site either continuously or in a switched (pulsed) manner. The main benefit of switching enables power conservation and facilities much higher  
15 peak power output, thereby improving clinical response.

Preferably, electromagnetic radiation therapy system also includes means for reducing the amount of ambient radiation which impinges on the site of infection. The presence of ultraviolet light and violet light as in sunlight exacerbates herpetic conditions and it is preferred to exclude wavelengths below 400nm. More  
20 preferably, wavelengths below 500nm are excluded.

Preferably the system further includes means for fixing the intensity of the radiation within a pre-determined range. The radiation output may be monitored with a visible  
25 display indicating correct function of the device both for intensity and wavelength.

Preferably the system further includes means for controlling the duration of the application of the radiation. Accordingly, the present invention is concerned with the use of electromagnetic radiation having a wavelength in the range from visible to  
30 infra red and applied at a low intensity such that no thermal damage is caused to any human or animal tissues.

In the case where the system is to be used in such a way that radiation will be caused to enter the eye, it is preferred that the power intensity does not exceed 100 mWatts/cm<sup>2</sup>. Otherwise, the power intensity may be higher and can suitably be 5 delivered in pulsed form, thereby obtaining several watts of momentary power output, allowing good penetration of tissue and substantial systemic effect.

The radiation producing means are preferably solid state light emitting devices, more preferably solid state light emitting diodes or gas discharge devices. The radiation 10 from such devices can be electrically operated or the radiation can be delivered to an applicator via a fibre-optic delivery system.

Preferably, the radiation emitter includes a PN junction arranged to emit radiation with a wavelength centring at or about 1072nm or at or about 1268 nm. A single 15 light diode assembly may include a plurality of orientated junctions.

Infrared emitting diodes may be arranged not only to emit radiation at a specific frequency but also to emit a high intensity divergent beam.

20 A gas discharge device may include a mixture of gases which will give an output at the desired wavelength, for instance, 1072 nm.

Another preferred radiation producing means is a laser diode device, an example being a laser diode emitting light at a frequency of 1064.nm. Such a light emitting 25 means is of low power intensity having a divergent beam and not giving rise to thermal damage. It may be used to treat many conditions, including pain relief.

The present invention also provides the use of divergent electromagnetic radiation having a wavelength of between 980 and 1500 nm and an intensity of at least 30 50μWatts/cm<sup>2</sup> and up to 2 Watts/cm<sup>2</sup> to treat an area of biological tissue of a living human or animal subject.

Preferably the electromagnetic radiation as produced by the system of the invention provides for treating conditions such as, without limitation, herpetic infections, bacterial and/or viral infections of the skin or upper respiratory tract, ophthalmic conditions such as "dry eye syndrome", caustic injuries, musculoskeletal conditions, inflammatory conditions such as rheumatoid arthritis and malignancies, reduction of scarring, promotion of wound healing, sports performance and providing acute and chronic pain relief.

- 10 The use of restricted bandwidth radiation can enhance the immune system as a result of which the body is able to combat infections, such as the herpes virus.

Although reference has been made to infections caused by the herpes virus, the present invention is not limited to such infections. It is applicable to other infections 15 caused by all viruses including HIV, common cold and influenza viruses.

- The present invention also provides a method of treating an area of biological tissue of a living human or animal subject comprising applying to said area divergent electromagnetic radiation having a wavelength of between 980 and 1500 nm at an 20 intensity of at least  $50\mu\text{Watts}/\text{cm}^2$  and up to  $2\text{ Watts}/\text{cm}^2$ .

Preferably, the area to be treated is irradiated so that the affected tissue receives at least  $50-500\ \mu\text{Watts}/\text{cm}^2$  peak power of radiant energy, depending on the tissue to be treated. A factor here is the period of irradiation and, preferably, the period should 25 be at least a specified minimum of  $10-15\ \mu\text{seconds}$  at a repetition rate /frequency of 450-800 Hz and preferably for at least 30 seconds duration.

#### Brief Description of the Drawings

- 30 Embodiments of the invention will now be described, by way of examples only, and with reference to the accompanying drawings, in which: -

Figures 1 to 4 are a view with cover removed, side view, under view and front view respectively of a first embodiment in accordance with the present invention;

Figures 5 to 7 are a front view, top view and under view of a second embodiment in accordance with the present invention;

5 Figures 8 to 10 are a back view, top view and a side view of a third embodiment of the present invention;

Figures 11 and 12 are a side view and a view from the right (as seen in Figure 11) of a fourth embodiment in accordance with the present invention; and

Figures 13 and 14 show further embodiments of the present invention.

10

### **Detailed Description of Preferred Embodiments**

Referring to Figures 1 to 4, a first embodiment in accordance with the present invention includes a hand held divergent narrow wavelength radiation source 4 with 15 a built in timer and ambient radiation detector. A single wavelength is used at any one time, preferably in the infrared spectrum. However, the effective wavelengths, which may be covered by such a device, extend from the visible spectrum to the infrared. In another embodiment of the invention, two wavelengths are used, one that is visible and the other that is invisible, particularly in the case where the optimal 20 wavelength is in the infrared.

Radiation source 4 includes an elongate, rectangular cross-section hollow body with one end 1 being transparent to light. The radiation source includes an array of light emitting diodes 2 mounted close to transparent end 1. Power is delivered to devices 25 2 by means of batteries 3 located within the body 4.

The radiation source is provided with two On/Off switches 5, which may be actuated to initiate the operation of the internal electronics. Both buttons 5 have to be pressed simultaneously in order to operate the device correctly, thereby preventing 30 inadvertent use of the device. Close to the end opposite transparent end 1 is a utility

hole 6 which allows the radiation source to be hung up or attached to another article such as a bunch of keys.

The radiation source is provided with control electronics, which limit the time that  
5 the radiation source is on and then automatically switches off the radiation source. The control electronics monitor the ambient radiation and, in the event that the ambient radiation is of an intensity that would interfere with the therapeutic effect of the radiation source, an alarm buzzer (not shown) sounds. The radiation emitting devices 2 and their location and arrangement within the radiation source are such that  
10 the radiation emitted from the radiation source is in the form of a divergent light beam. Flange 8 restricts the ambient radiation incident on the area whilst being treated.

The radiation therapy system of the present invention could be adapted to be portable  
15 and for use by those individuals wishing to have their own self-contained and battery operated devices. Additionally it could be adapted so that during use an animal or human appendage could be located within a hollow body 3 to receive radiation emitted therefrom. This particular modification is particularly suited to treatment of genital herpes around the scrotal sac and/or penis.

20 Referring to Figures 5 to 7 of the accompanying drawings, a second embodiment in accordance with the present invention is in the form of a multi-panel narrow wavelength radiation source. In this case, a plurality of panels 3 are mounted in a side by side relationship on hinges 7 which, in turn, are connected to a stand 9 by  
25 means of arms 8 and 10. The arrangement is such that the panels can move relative to each other and the stand can be adjusted to alter the direction of illumination. The stand either extends from the floor or is attached to a chair or bed.

30 The front wall of each panel 3 is transparent and, mounted below the front wall, is an array of radiation emitting devices 4.

As with the earlier above described embodiments this embodiment of the invention includes control electronics to limit the time of the application of the radiation and to monitor the ambient radiation and provide an alarm when the threshold value of the ambient radiation is exceeded.

5

Referring to Figures 8 to 10 of the drawings, a third embodiment in accordance with the present invention is in the form of a narrow wavelength radiation source with adjustable headgear.

- 10 The radiation source is, in use, located on the operator's head and includes two panels 1, 5 of radiation emitting devices, panels 1 being separated by an intervening notch 1a. These radiation panels 1 can be used either simultaneously or separately, there being provided a switch (not shown) to direct electrical power to one or both of panels 1.

15

The radiation panels 1 are held close to the eyes by adjustable control elements 2.

- 20 The radiation source is provided with control electronics 4, which limit the time of application of the photons to the affected site and also automatically switch off the radiation at the end of the application period. As before the control electronics monitor the ambient radiation and provide an alarm when the threshold level is exceeded.

- 25 Referring to Figures 11 and 12 of the accompanying drawings, a fourth embodiment of the present invention is in the form of a narrow or restricted bandwidth radiation source for delivery of photons to an orifice. In this case the body of the radiation source includes an elongate cylindrical portion 2 having at one end a flange 4 whose shape is indicated in Figures 11 and 12. At its other end, elongate portion 2 is hemispherical. Radiation emitting devices are located both in the elongate portion 2 and the flange 4 and this radiation source can be used to deliver photons to any orifice in the human/animal body, for instance, the vagina, anus, oro and

nasopharynx and buccal cavity. The radiation source may be provided in different sizes according to the size of the orifice into which it is to be inserted.

- Control electronics limit the time of irradiation and monitor the ambient radiation, as  
5 with the previously described embodiments of the invention.

Figures 13 and 14 illustrate devices useful in the treatment of the common cold and acne.

- 10 The common cold is caused by a viral infection of the upper respiratory tract. The viral particles are almost exclusively found in the pharynx, sinuses and nasal passages.

15 The device is a radiation emitting apparatus, which delivers a narrow bandwidth radiation, which, is of a wavelength, that will penetrate the superficial skin and penetrate the underlying tissue to sufficient extent to generate a therapeutic effect.

The device in Fig 14 is flexible and is placed against the patient's face whilst he is lying supine. Notch 1 provides an aperture for the patient's eyes. Panel 2 provides  
20 treatment for the frontal sinuses. Panel 3 provides treatment for the maxillary sinuses and nose, and the intervening bridge 1a provides treatment for ethmoid sinuses and nose.

25 The device in Fig 13 is a flexible radiation emitting apparatus, which is placed against the patient's neck so that the points 5 approximate the base of the patient's ears. This apparatus delivers radiation for therapeutic effect to the patients larynx, oropharynx and laryngopharynx. Depression 4 fits underneath the patient's chin.

Devices for treating acne are as shown in Figures 13 and 14 and comprise several  
30 panels of variable shape and size. All the panels have a radiation emitting surface 2 and are flexible to enable the panel to follow the contour of the face and neck. Panels

6 and 7 (Figure 13) are applied to the inferior aspect of the chin and the neck respectively.

5 Panels in the Figure 14 device are applied to the face so that notch 1 enables the patient to see whilst being treated. Bridge 1a treats the bridge of the nose and extension 3 the cheeks. Notch 4 rests on the tip of the nose allowing the patient to breathe comfortably during treatment. Panel 8 is used to treat the chin area and the area adjacent to the mouth.

10 Where the chest and/or back is involved a larger version of panel 6 would be used.

Due to the superficial nature of the pathology ambient radiation is of significance and an ambient radiation detector is utilised.

15 Treatment time is at most 10 minutes but typically 5 minutes.

It should be appreciated that two or more of the above-described devices can be used in conjunction with one another. An example is in the treatment of paediatric herpetic stomatitis where the device for delivery of radiation to an orifice could be 20 used in conjunction with the multipanel device to ensure adequate delivery of the radiation system.

The application of radiation in the manner described appears to give the patient immediate (within 6 minutes) relief from any pain which is chemically mediated 25 irrespective of the inflammatory condition causing it i.e. it gives pain relief in conditions other than those caused by viral infections. It does not affect the conduction of pain impulses as in local anaesthetics.

### **Experimental Results**

30 Examples of the invention will now be described with reference to the treatment of particular conditions.

**Herpes**

The radiation-emitting surface was placed against the cold sore for at least 90 seconds, typically 4 minutes. The environmental conditions must be such that the ambient radiation to the cold sore must be decreased below acceptable levels. This can be achieved either by switching the radiation off or designing the device with a flange around the outside, diminishing the ultraviolet light to the area. Treatment is only once a day. One treatment may be all that is necessary however, in view of the fact that this also enhances wound regeneration and the wound regenerative effect only lasts 24 hours, daily treatments would improve clinical response. Various wavelengths were evaluated using a double blind control trial, Zovirax being given to the control patients. The average time for patient to be treated with 660 nm radiation took 7.5 days. The average time for a patient to be treated with 1072nm radiation was 3 days if lesion was already present. However it was less than 12 hours if the patient only has a tingling sensation.

99.5% of patients abort their attack if they are treated within the tingling period using 1072nm radiation. The cure rate of the patients receiving radiation treatment was total, in that none had a recurrence of their cold sores at the site treated. However 20% of the acyclovir treated group had recurrence at the site of treatment.

The number in the trial was 300.

A further study was conducted in a double blind protocol comparing radiation treatment to acyclovir. The group receiving radiation therapy had cold sores that healed within 4.7 days and the group receiving acyclovir had cold sores that healed within 4.7 days. Statistical analysis resulted in a p value of 0.027, which was statistically significant.

Please refer to the table below for further details.

Treatment	Number of Patients	Mean Time to Healing (days)
Placebo radiation plus acyclovir	14	6.9
Active radiation	15	4.7
Active radiation plus placebo cream	16	6.7
Acyclovir	18	8.5
Red Light	20	7.5

### Genital herpes

- Again the applicator was required to follow the contours of the genitals and for a woman the cervix and posterior fornix was treated simultaneously with vagina and perineum. The treatment period is only 4 minutes. Due to the shape of the device, ambient radiation is excluded from the treatment area, and darkened room is not necessary.
- 10 A total of eight patients have been treated, all of who have reported a shorter duration of their recurrent attacks of genital herpes. In addition, after one year of treatment, for each attack, all patients have been free from recurrence for the last 6 months, whereas they typically had an attack per month.

### 15 Shingles

- The radiation panel was applied to the area that is infected, for periods of 4 minutes. If the panel is applied to the actual skin surface and the panel is optically opaque it will occlude the ambient radiation from the area and hence allow treatment to proceed successfully. The only exception where a darkened room is essential is for the treatment of ophthalmic conditions when one cannot have the radiation-emitting surface close to the eye because of the generation of heat. The generation of heat in association with ophthalmic conditions is contraindicated. A helmet was used with an ambient radiation detector with alarm so that the radiation can be delivered to the

orbital region. Again the treatment period was 4 minutes. Generally ophthalmic conditions are treated on a daily basis. However, ophthalmic herpes can be treated once every three days to achieve a positive result.

- 5 Four patients have been treated; all of who had significantly shortened healing periods and decreased incidence or recurrence.

#### **The common cold**

The patient was reclined on a bed, preferably in subdued lighting, but this is not  
10 essential. The device was placed against the skin ensuring that the frontal, ethmoidal and maxillary sinuses are covered. Another panel was placed again the neck, ensuring that the device goes as high as the angle of the jaw to enable treatment of the pharynx and larynx areas. The treatment cycle is at least 4 minutes. After that has been completed an oral device was used which is very similar to the vaginal  
15 applicator without the flange. The device was inserted into the patient's mouth. The treatment period was again for 4 minutes. This applicator has a disposable outer skin, which is changed with each patient. The applicator will treat the soft palate and the back of the oral pharynx and the top of the nasopharynx where the surface applicator would not achieve acceptable penetration levels. Using this protocol  
20 alleviation of the pain associated with pharyngitis was achieved immediately, i.e. within 90 seconds, and the symptoms associated with oral pharyngitis have been alleviated within six hours.

Ten patients have been treated. Of note is that unrestricted light can be applied to the  
25 pharynx with immediate relief of symptoms, however for improved efficacy, light of the wavelength 1072nm is required to be applied to the face and sinuses. We noted that in all cases the sore throat was improved immediately and symptoms of congestion and flu-like illness were alleviated in 4-6 hours.

**Acne**

Using the same applicator adolescent acne can be treated by the simple addition of a chin extension. The treatment time is 4 minutes. Ambient light is important but not in as much as the applicator will have an opaque surface so therefore it will be sheer  
5 proximity to the skin reduce ambient light to the skin. Applications should initially be every two to three days and maintenance would be perhaps once a week.

Twelve patients have successfully been treated with light of the light therapy system.  
Results indicate that if 950nm .5mm diodes were used there was a marked  
10 exacerbation of the disease process, however when using 950nm 8mm diodes there was a slight increase in inflammation but an overall improvement. The use of restricted radiation resulted in a rapid resolution of acne over 7 days. Daily treatment enhanced clinical results. Once a clinical result was achieved, the regimen was maintained for 1-2 treatments per week for a satisfactory outcome.

15

**Musculoskeletal disorders**

Treatment of musculoskeletal disorders such as tennis elbow gout, muscle injuries and knee injuries. The application time again is only 4 minutes. The radiation is directed over the affected area and gentle pressure is applied. Once the treatment is complete the patient feels immediate pain relief and improved joint movement. Muscle stiffness is greatly improved. This is a distinct advantage in the treatment of gout since this can be an extremely painful condition. The treatment could be repeated after 24 hours. Generally speaking treating prior the 24 hours is not indicated, as there is no additional clinical benefit.

25

**Post-operative wounds**

24 hours after an operation, a 4-minute treatment period reduces pain for 6-8 hours and this was repeated three to four times a day to enhance wound healing. The treatment may be used on a daily basis with or without the pain for post-operative  
30 wounds.

Seven volunteers, all of whom had minor surgery found decreased scar formation if the area of surgical incision was treated daily for 10 days commencing on the day of surgery. Thus it is envisaged that the present invention has applications in the treatment of keloids, burns and cosmetic surgery.

5

### **Connective Tissue Diseases**

Rheumatoid arthritis is an example of this group of conditions. The painful areas are treated in a device which comprises one fixed panel in which the hands are placed on and a flexi panel which is placed over the top of the hands applying gentle pressure 10 to aid additional penetration of the skin. The treatment time is 4 minutes. Ambient radiation does not appear to be a significant factor in the treatment of rheumatoid arthritis. Part of the treatment protocol can also be in the treatment of thymus, liver and regional lymph nodes, which are all associated with antigen recognition. Again treatment time is 4 minutes. The thymus, lymph nodes, liver and spleen may be 15 treated once a week, whereas the hands may be treated initially once a week. However, during an acute exacerbation they can be treated daily. If treated more than once daily there appears to be no advantage.

A small trial involving eight patients in the Rheumatology Clinic resulted in the 20 findings that those receiving placebo radiation treatment felt more relief than those patients receiving active 950nm radiation. However, when using radiation in the restricted wavelength according to the present invention, all ten patients reported a clinical benefit compared to the placebo treatment. In addition we found that if the thymus and other aspects of the reticuloendothelial system were treated with the 25 restricted radiation the patients experienced an overall cessation of their arthralgia and myalgia.

### **Malignancy**

The area of the tumour is treated together with treatment to the antigen recognition 30 centres such as thymus, spleen, liver and lymph nodes. Daily whole body treatments may be carried out.

**Bacterial Infections**

A number of diverse minor bacterial infections have been treated successfully in 35 patients. In all cases the infection had been present for at least three days and was considered to be worsening daily. At the time of treatment each patient would have ordinarily been prescribed an oral antibiotic. In 70% of cases the infection was resolving 6 hours after treatment and was completely better within one day. In the remaining 30% of cases the infection had improved considerably within one day and following a second radiation treatment, the infection was gone by the second day. Of clinical note was the observation that in the Caucasians treated there was a significant decrease in superficial scar tissue at the site of assault. As previously noted, the 8mm 950nm diode unrestricted radiation source was effective but not as effective as the restricted radiation of 1072nm of the present invention.

**15 Sports Medicine**

Treating all muscle groups prior to training will increase the level to which the athlete can train by as much as 50%, in addition to decreasing the incidence of muscle injury.

- 20 In a double blind trial, five volunteers were used by treating one limb with placebo radiation and the corresponding other limb with active radiation. In all cases the volunteers were able to increase their effort tolerance before feeling muscle fatigue by 30-50% in the limb treated with the active radiation.
- 25 Muscle injuries have been successfully treated in 35 patients. The pathology involved included rotor cuff syndrome, tennis elbow, lower back pain and lumbar fascitis. The response to therapy ranged from immediate relief in 30% of cases to complete relief after 24 hours in the remaining 70%. Daily treatment is required until the problem is resolved.

### Ophthalmic Conditions

Chronic ophthalmic pain was successfully alleviated permanently in 90% of the eleven patients treated.

5 Unstable corneal epithelium (over a period of 6 weeks) which resulted in recurrent corneal ulceration, was stabilised in 6 patients promoting an intact corneal surface within 5 days of commencement of daily treatments of radiation therapy.

10 Four patients with conjunctivitis sicca or dry eye syndrome, reported significantly decreased eye irritation and produced significantly less debris accumulation within fornices following radiation treatment. Of note was the experience that whilst radiation centred on 1072nm was effective in this condition, radiation centred on 1268nm was more effective. Once weekly treatment with radiation was sufficient to alleviate symptoms.

15 Episcleritis and other inflammatory conditions of the eye were successfully treated in 11 patients. Daily treatments were necessary to obtain the desired clinical effect. The recurrence of the inflammatory condition was decreased in all cases. All patients treated had suffered their conditions over several months and resolution occurred in 20 3-4 days without the use of eye drops.

25 Severe caustic injury to the eye is considered untreatable and almost always results in destruction of the cornea and blindness. Animal experiments (conducted in South Africa) have indicated that caustic injury is treatable with the radiation of the present invention.

30 Five pairs of rabbit eyes were exposed to a supersaturated solution of NaOH for 30 seconds after topical administration of an anaesthetic. All eyes were washed out thoroughly following the caustic injury and it was noted that the corneas were immediately opaque following the injury. One eye of each rabbit was then either treated with conventional steroids plus antibiotic whilst the other eye was treated

with the radiation of the present invention. All rabbits were sacrificed 4 weeks following twice daily treatment. In all cases the eyes that had been treated with conventional therapy (steroids plus antibiotic) rapidly developed panophthalmitis with resulting blindness, whilst the eyes treated with the radiation therapy of the 5 invention showed that the anterior chamber, lens, vitreous and retina were intact despite corneal damage. In addition the scar tissue in the radiation treated eyes was reduced by at least 50% compared to the conventionally treated eyes.

Further experiments involving a less concentrated solution of NaOH being applied 10 for a longer duration, 3 minutes, resulted in corneal damage to all eyes as gauged by pale milky appearance to the corneas. The same protocol of treatment was applied, I.E. one eye being treated in conventional manner and the other by radiation therapy. The animal were sacrificed following two weeks of treatment and the results showed that eyes treated with the radiation therapy were clear whilst the conventionally 15 treated eyes remained opaque.

The ability to reduce scarring was investigated using 5 pairs of rabbit eyes. Each eye was scarred by a 4mm linear full thickness incision in the centre of the corneas under topical anaesthesia. One eye was treated conventionally and the other by radiation 20 therapy. All eyes healed however the eyes receiving radiation therapy healed at a faster rate with scarring reduced by 50%. It was noted that radiation centred on 1072nm gave best similar results.

### Pain

25 Radiation at 950 nm was only marginally effective compared to the restricted radiation at 1072 and 1268 nm. It was noted that 1072 nm was more effective than 1268 nm radiation in treatment of acute pain as caused by a superficial burn. However, 1268 nm radiation was reported as more effective at alleviating deep 30 muscle pain caused by muscle injury.

Pain was gauged in patients by subjective assessment.

Thus it will be appreciated that the radiation therapy system of the present invention has wide application in treating a variety of different diseases and conditions. The radiation therapy system effects on a patient are rapidly felt and since the system is non-invasive it presents a less stressful/traumatic therapy to the patient. Moreover,  
5 the system could be used to treat a wide variety of patients quickly thus reducing the financial burden to the health service.

**AMENDED CLAIMS**

1. An electromagnetic radiation therapy system comprising means for emitting divergent electromagnetic radiation between 980 nm and 1500 nm and being capable of producing, at the site being treated, a radiation intensity of at least 50  $\mu$ Watts/cm<sup>2</sup> and up to 2 Watts/cm<sup>2</sup>.
2. An electromagnetic radiation therapy system according to either of Claims 1 or 2 wherein the wavelength is in the range 980nm-1300nm.
3. An electromagnetic radiation therapy system according to any preceding claim wherein the wavelength is at, or about, 1072nm.
4. An electromagnetic radiation therapy system according to any preceding claim wherein the wavelength is at, or about, 1268nm.
5. An electromagnetic radiation therapy system according to any preceding claim wherein the half angle divergence of the electromagnetic radiation is in the range 15° to 45°.
6. An electromagnetic radiation therapy system according to any preceding claim wherein the electromagnetic radiation is continuous or pulsed.
7. An electromagnetic radiation therapy system according to any preceding claim wherein, in the instance of the electromagnetic radiation being continuous, the intensity is at least 50  $\mu$ Watts/cm<sup>2</sup> for treatment of eyes and mucous membranes and up to 2 Watts/cm<sup>2</sup>.
8. An electromagnetic radiation therapy system according to any preceding claim wherein, in the instance of the electromagnetic radiation being continuous, the intensity is at least 500  $\mu$ Watts/cm<sup>2</sup> for treatment of skin and up to 2 Watts/cm<sup>2</sup>.

9. An electromagnetic radiation therapy system according to any of Claims 1-6, wherein in the instance of the electromagnetic radiation being pulsed, the intensity is at least 50  $\mu$ Watts/cm<sup>2</sup> peak power for treatment of eyes and mucous membranes and the average power is up to 2 Watts/cm<sup>2</sup>.
- 5
10. An electromagnetic radiation therapy system according to any of Claims 1-6, wherein in the instance of the electromagnetic radiation being pulsed, the intensity is at least 500  $\mu$ Watts/cm<sup>2</sup> peak power for treatment of skin and the average power is up to 2 Watts/cm<sup>2</sup>.
- 10
11. An electromagnetic radiation therapy system according to any of Claims 1-6 or 9 or 10 wherein the average power of the pulsed electromagnetic radiation intensity is in the region of 50-100 $\mu$ Watts/ cm<sup>2</sup>.
- 15 12. An electromagnetic radiation therapy system according to any of Claims 1-7 or 9-11 wherein pulsed electromagnetic radiation is applied for periods of at least 10-15  $\mu$ seconds.
13. An electromagnetic radiation therapy system according to any of Claims 1-7 or 9-20 12 wherein the pulsed electromagnetic radiation is applied at a frequency/repetition rate in the range 480-800 Hz.
- 25
14. An electromagnetic radiation therapy system according to Claim 13 wherein the frequency/repetition rate is at, or about, 600 Hz.
15. An electromagnetic radiation therapy system according to any of Claims 1-7 or 9-14 wherein the pulsed electromagnetic radiation is applied to the affected area for at least 30 seconds and up to 15 minutes.
- 30 16. An electromagnetic radiation therapy system according to any preceding claim wherein the electromagnetic radiation therapy system also includes means for

## ART 34 AMDT

reducing the amount of ambient radiation which impinges on the site of treatment.

17. An electromagnetic radiation therapy system according to Claim 16 wherein the  
5 means for excluding ambient radiation excludes radiation below 400-500 nm.
18. An electromagnetic radiation therapy system according to any preceding claim  
further including means for fixing the intensity of the radiation within a pre-  
determined range.  
10
19. An electromagnetic radiation therapy system according to any preceding claim  
wherein radiation output is monitored with a visible display indicating correct  
function of the device both for intensity and wavelength.
- 15 20. An electromagnetic radiation therapy system according to any preceding claim  
further including further including means for controlling the duration of the  
application of the radiation.
21. An electromagnetic radiation therapy system according to any preceding claim  
20 wherein the radiation producing means are solid state light emitting devices,
22. An electromagnetic radiation therapy system according to Claim 21 wherein the  
solid state light emitting devices are solid state light emitting diodes or gas  
discharge devices or a laser diode device.  
25
23. An electromagnetic radiation therapy system according to either Claim 21 or 22  
wherein radiation from such devices is electrically operated or delivered to an  
applicator via a fibre-optic delivery system.

**ART 34 AMDT**

24. An electromagnetic radiation therapy system according to any of Claims 21-23 wherein the radiation emitter includes a PN junction arranged to emit radiation with a wavelength centring at, or about, 1072nm or at, or about, 1268 nm.
- 5      25. An electromagnetic radiation therapy system according to Claims 24 wherein a single light diode assembly include a plurality of orientated junctions.
- 10     26. An electromagnetic radiation therapy system according to Claims 22 wherein the gas discharge device may include a mixture of gases which will give an output at the desired wavelength, for instance, 1072 nm or 1268 nm.
- 15     27. Divergent electromagnetic radiation having a wavelength of between 980 and 1500 nm and an intensity of at least 50 $\mu$ Watts/cm<sup>2</sup> and up to 2 Watts/cm<sup>2</sup> for use in treating an area of biological tissue of a living human or animal subject.
- 20     28. Divergent electromagnetic radiation according to Claim 27 for treating herpetic infections, bacterial and/or viral infections of the skin or upper respiratory tract, ophthalmic conditions such as "dry eye syndrome", caustic injuries, musculoskeletal conditions, inflammatory conditions such as rheumatoid arthritis and malignancies, reduction of scarring, promotion of wound healing, improving sports performance and providing acute and chronic pain relief.
- 25     29. Divergent electromagnetic radiation according to Claim 27 for use in treating the immune system as a result of which a human or animal subject is able to combat infections, such as the herpes virus.
- 30     30. A method of treating an area of biological tissue of a living human or animal subject comprising applying to said area divergent electromagnetic radiation having a wavelength of between 980 and 1500 nm at an intensity of at least 50 $\mu$ Watts/cm<sup>2</sup> and up to 2 Watts/cm<sup>2</sup>.

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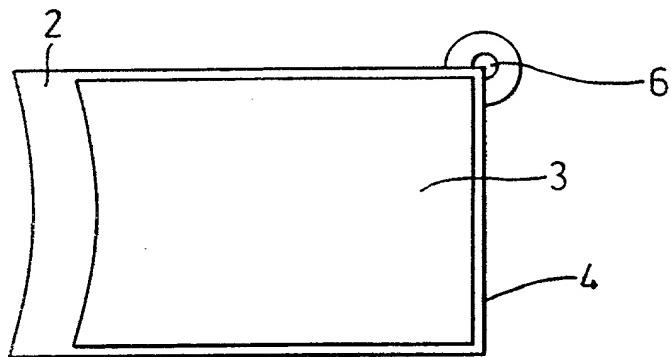


Fig. 1

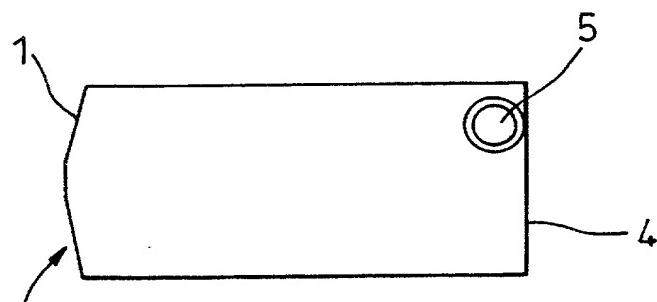


Fig. 2

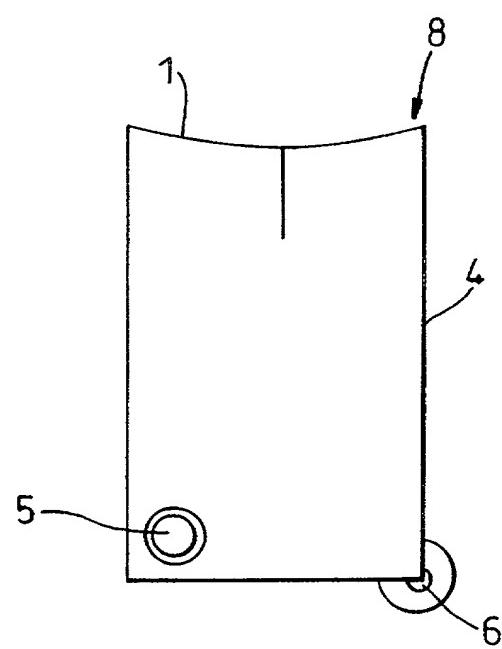


Fig. 3

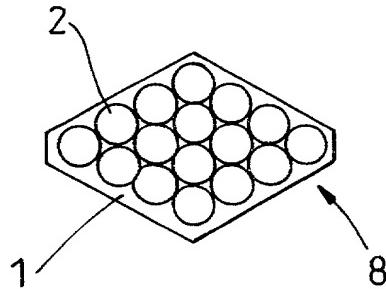


Fig. 4

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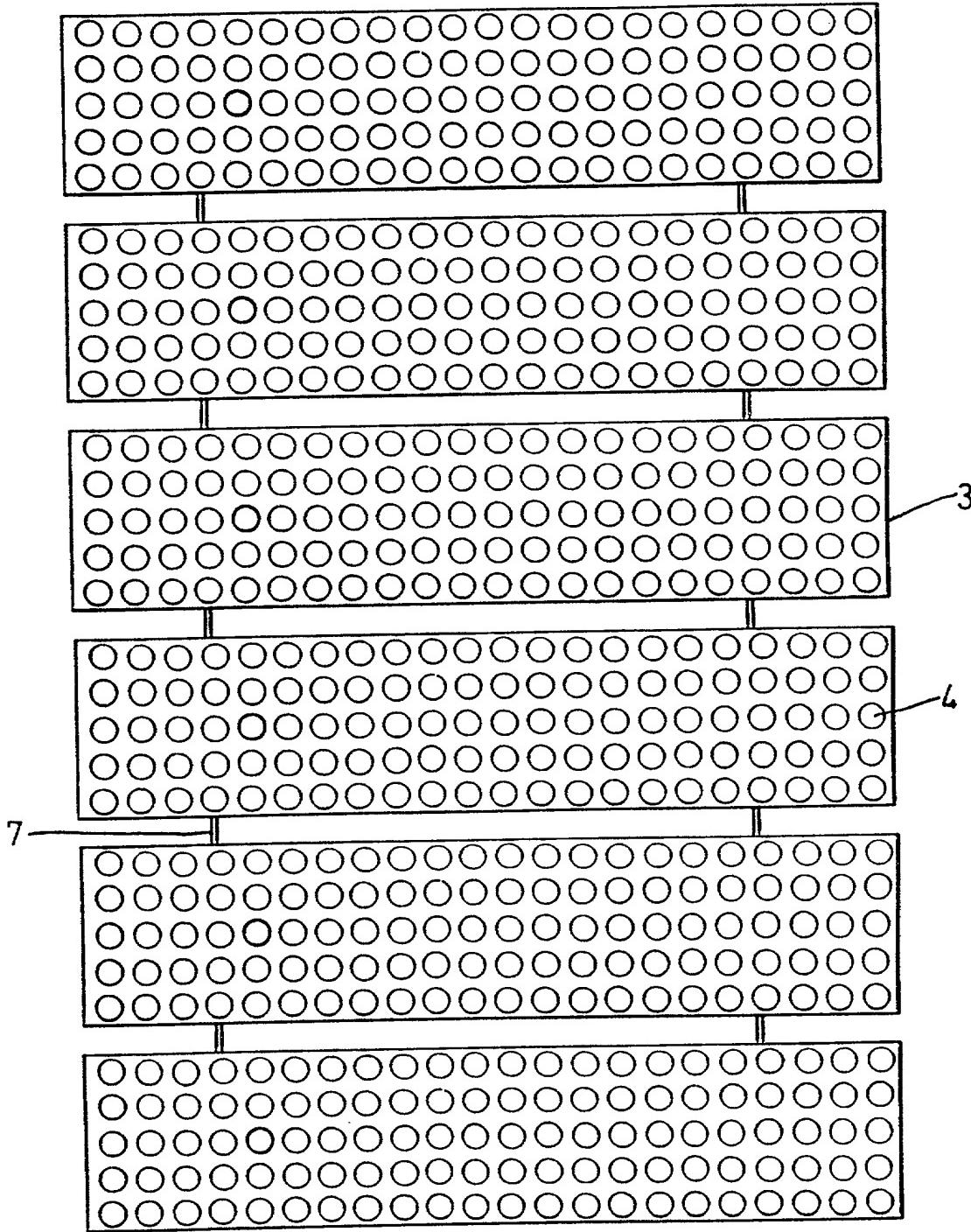


Fig. 5

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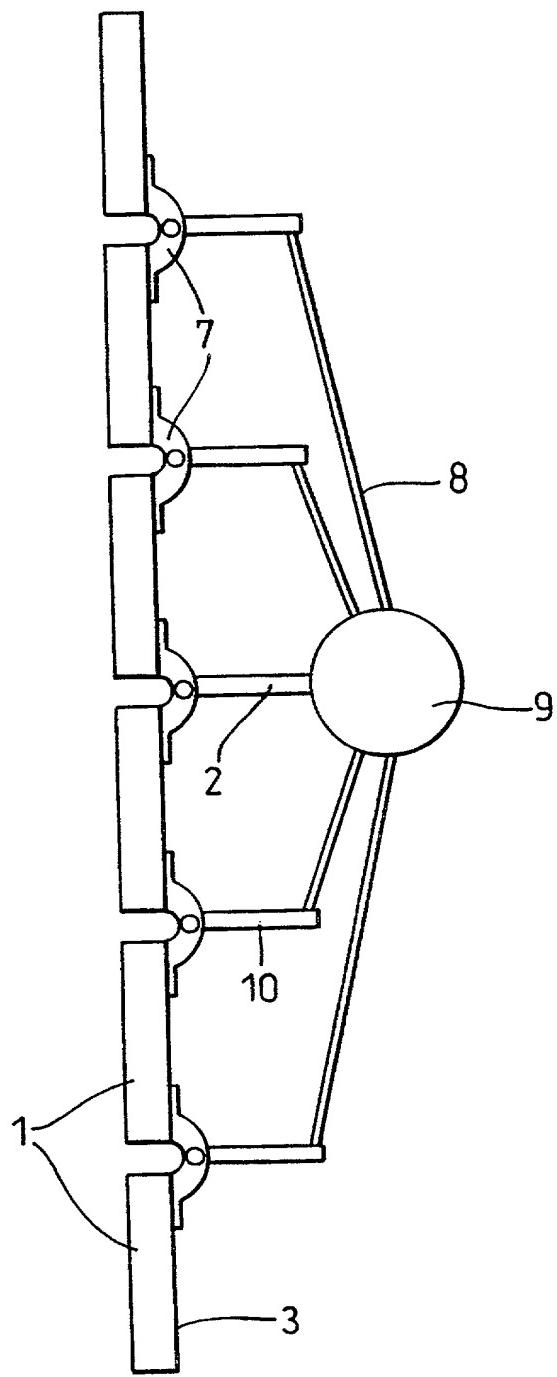


Fig. 6

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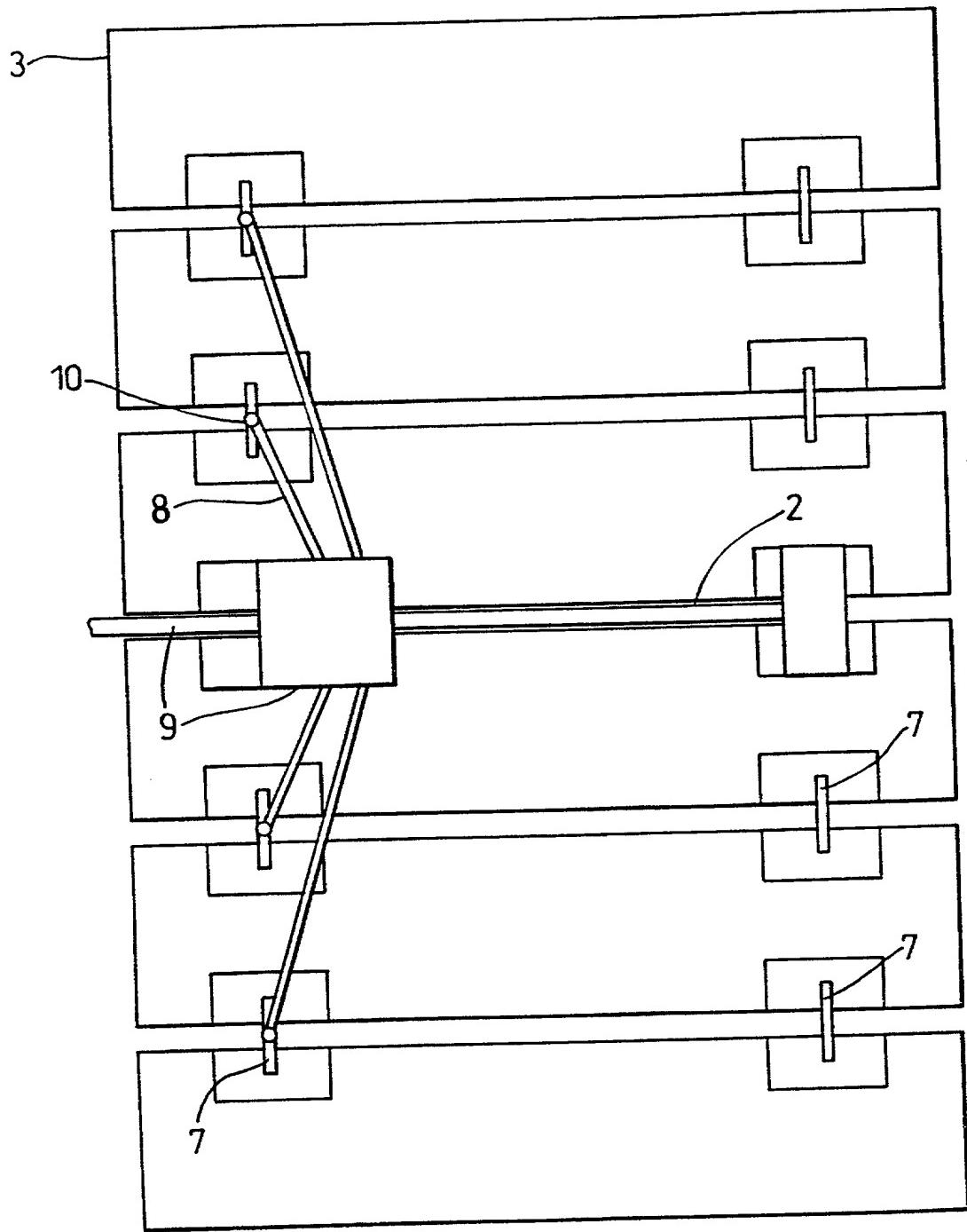


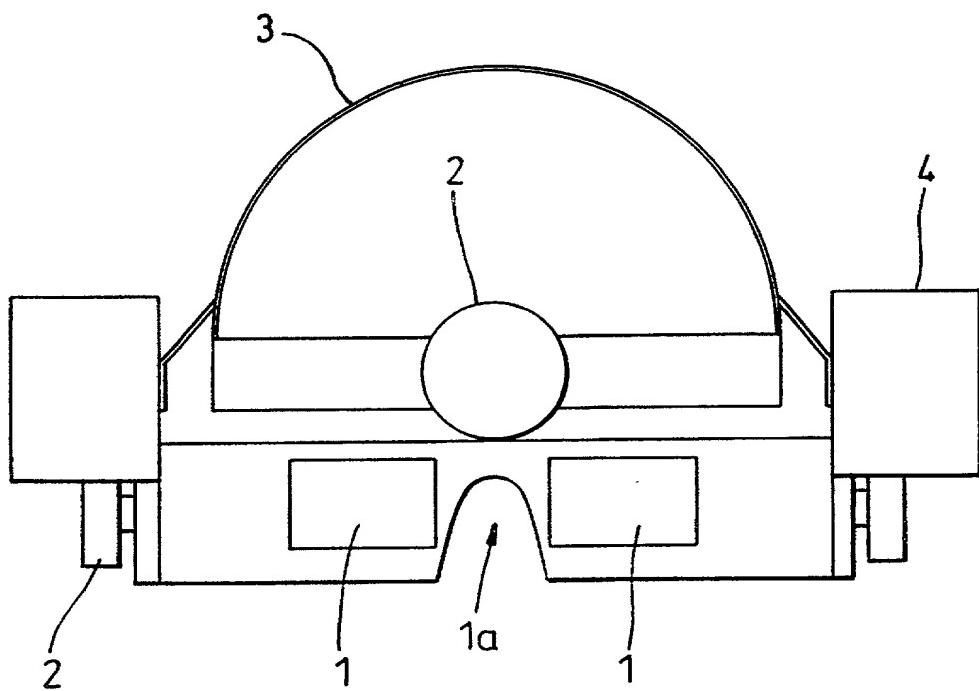
Fig. 7

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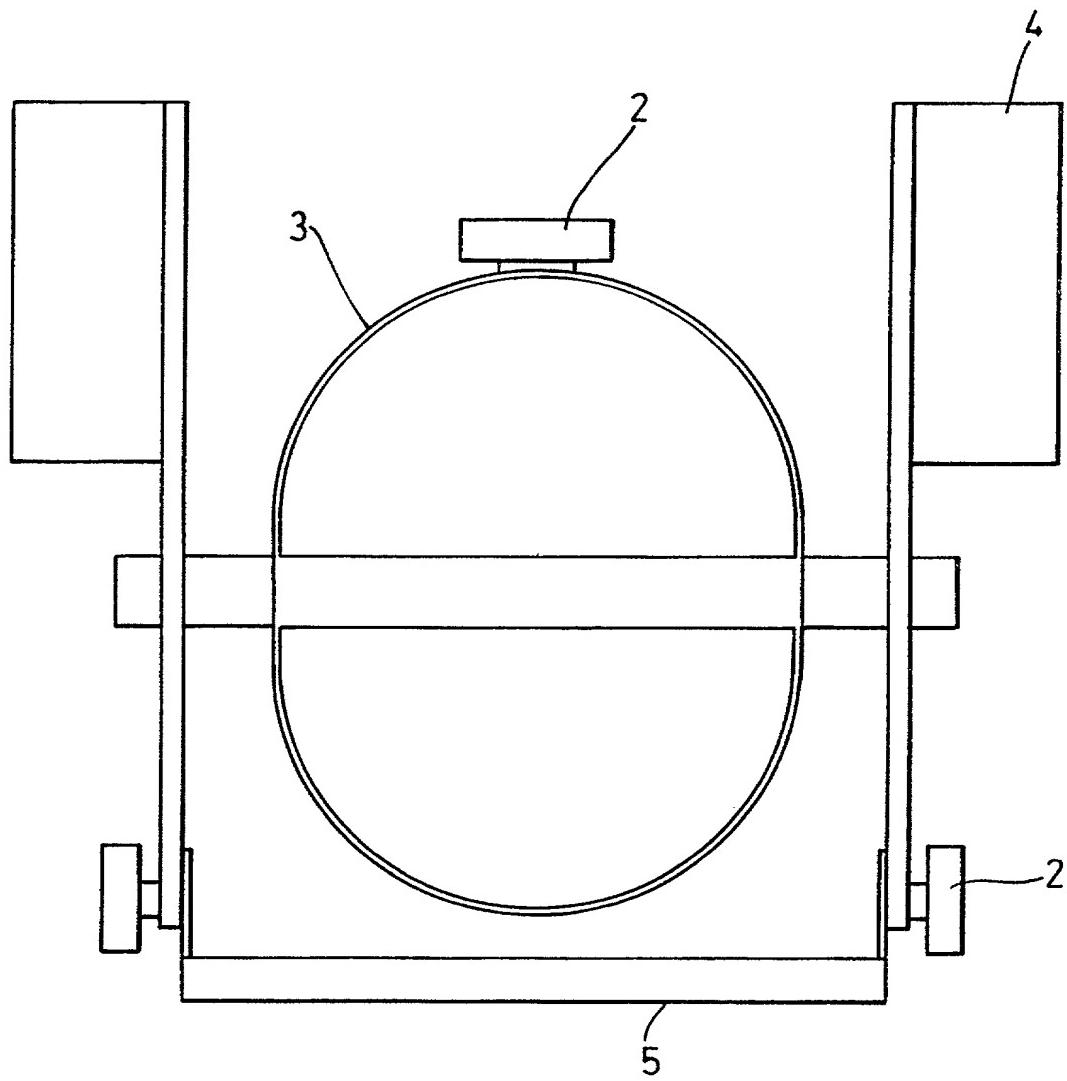
*Fig. 8*

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*Fig. 9*

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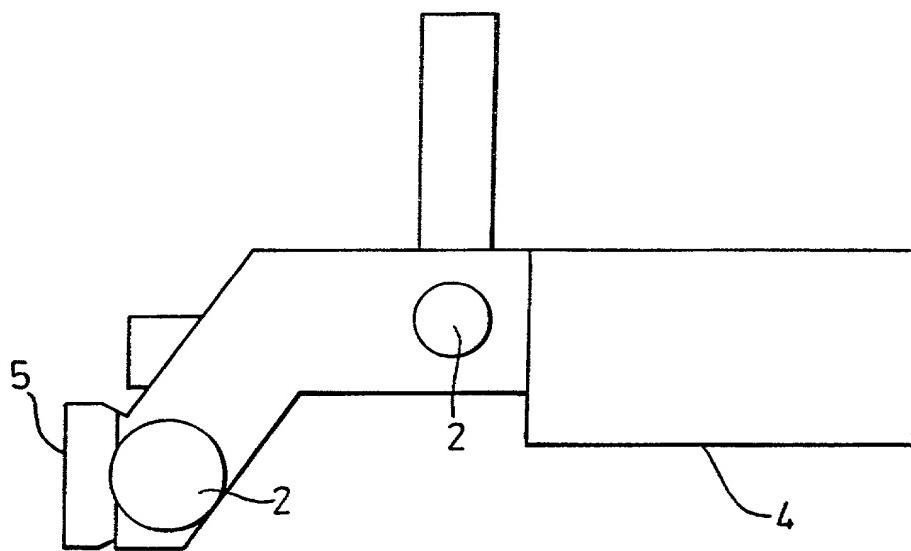


Fig. 10

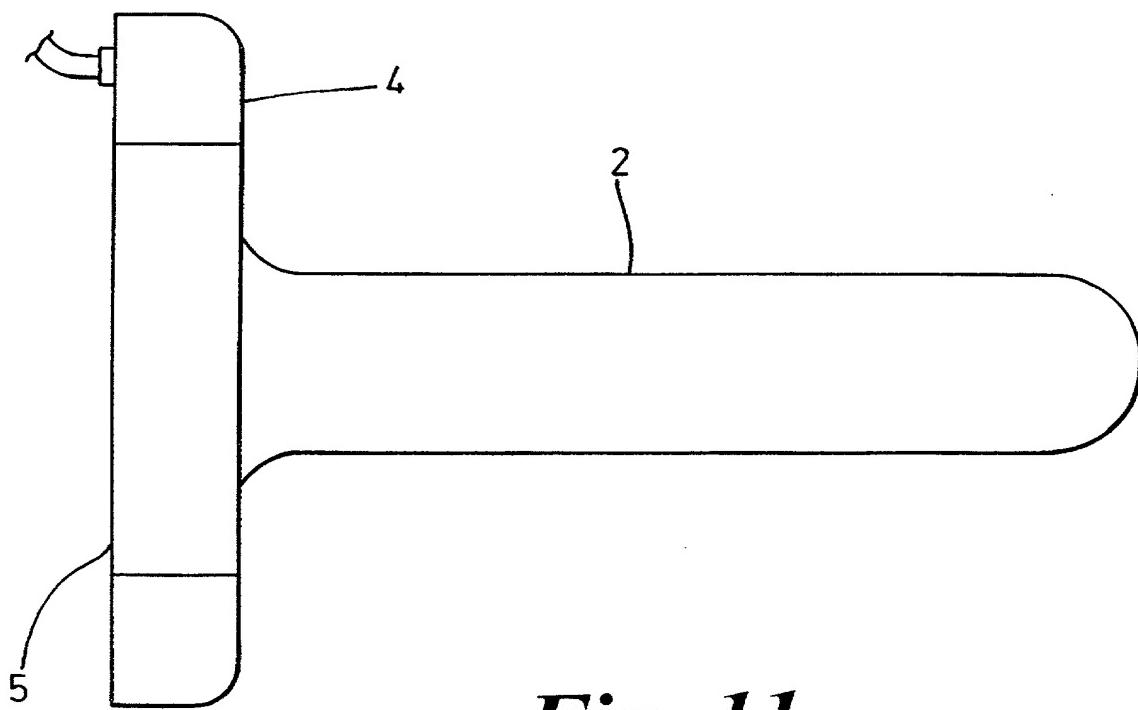


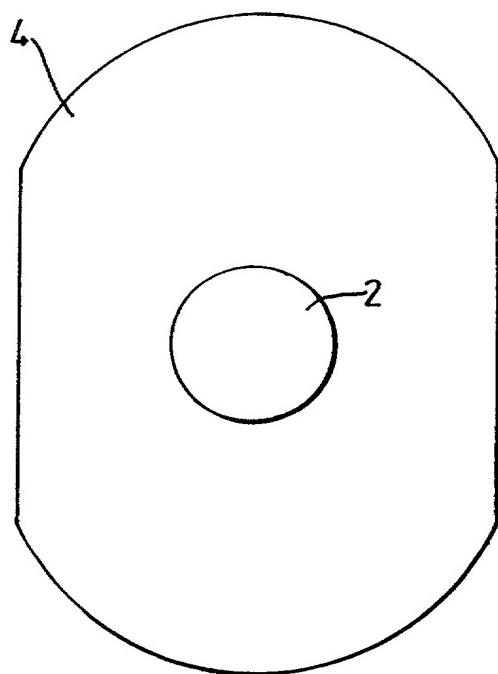
Fig. 11

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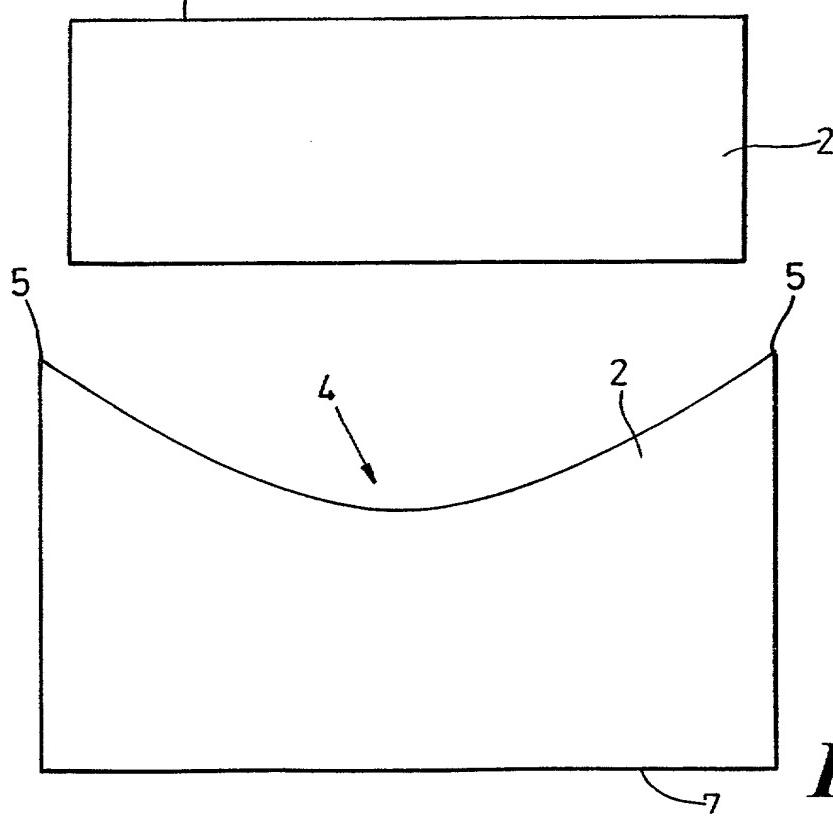
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*Fig. 12*



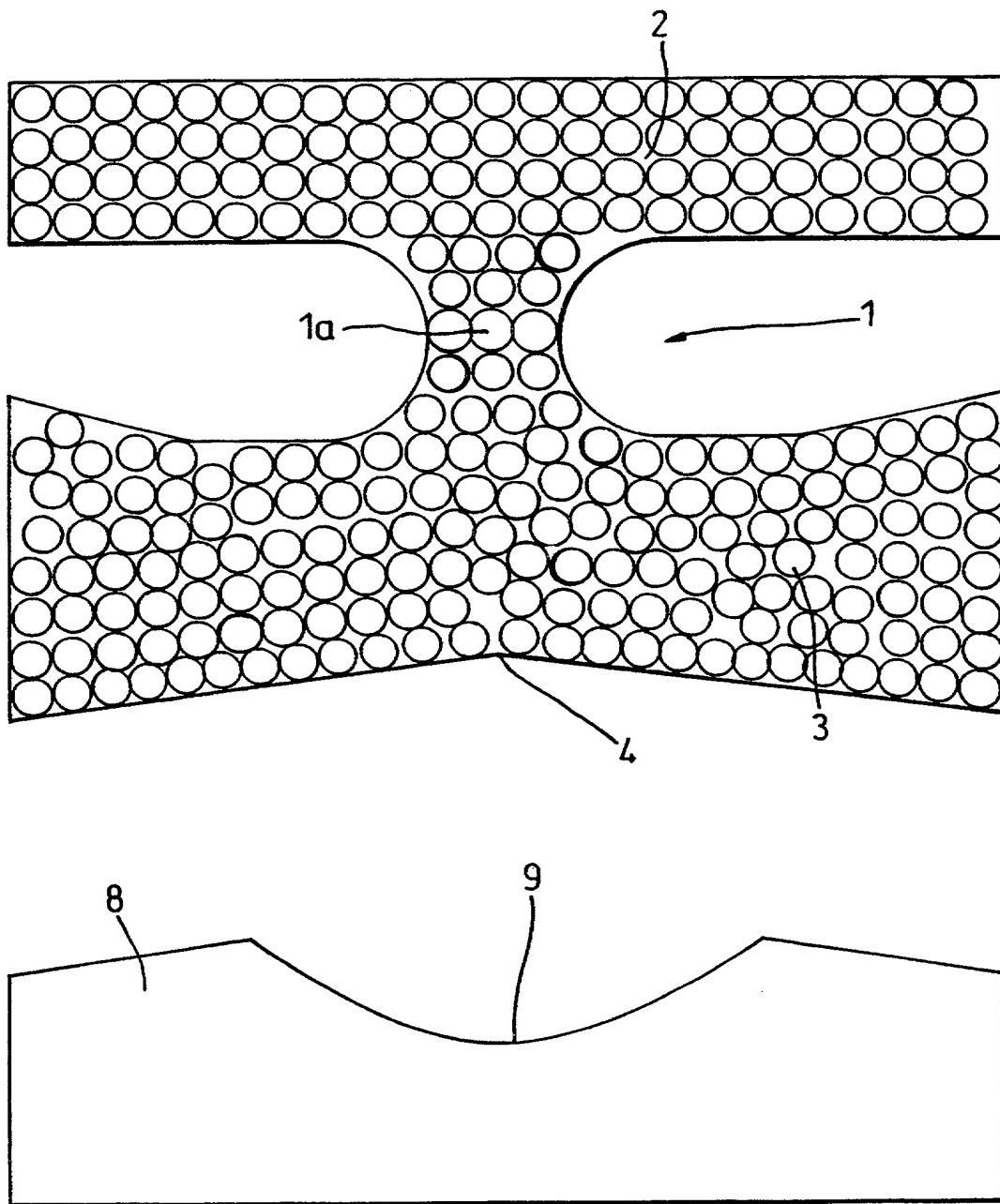
*Fig. 13*

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*Fig. 14*

**DECLARATION AND POWER OF ATTORNEY FOR PATENT APPLICATION**  
Attorney Docket No. 9052-53

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled **ELECTROMAGNETIC RADIATION THERAPY, ✓**

the specification of which

is attached hereto

OR

was filed on October 12, 1998 as United States Application No. or PCT International Application Number PCT/GB98/03073 and was amended on \_\_\_\_\_ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37 Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States of America, listed below and have also identified below any foreign application for patent or inventor's certificate, or of any PCT International application having a filing date before that of the application on which priority is claimed.

9721506.5	GB	10/10/1997 ✓	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
Number	Country	MM/DD/YYYY Filed	Priority Claimed
9727441.9	GB	12/31/1997 ✓	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
Number	Country	MM/DD/YYYY Filed	Priority Claimed

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below.

None	
Application Number(s)	Filing Date (MM/DD/YYYY)

Application Number(s)	Filing Date (MM/DD/YYYY)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) or § 365(c) of any PCT international application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application(s) in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application (37 C.F.R. § 1.63(d)).

None		
Appln. Serial No.	Filing Date	Status Patented/Pending/Abandoned

Appln. Serial No.	Filing Date	Status Patented/Pending/Abandoned
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Appln. Serial No.	Filing Date	Status Patented/Pending/Abandoned
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following registered attorney(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith.

**Customer Number 20792**

Send correspondence to:

Robert J. Smith  
Myers Bigel Sibley & Sajovec  
Post Office Box 37428  
Raleigh, NC 27627

Direct telephone calls to:

Robert J. Smith  
(919)854-1400

Facsimile:

(919) 854-1401

Full name of sole inventor: Gordon Rex Paterson Dougal

Inventor's

Signature: 

Date: 6/6/90

Residence:

Durham, Great Britain GBN

Citizenship:

British

Post Office Address:

6 Whitesmocks  
Durham  
DH1 4HW  
Great Britain